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CLAIMS:

- 1. A drug selected from the group consisting of at least one bisphosphonate when used for promoting new bone formation at a fracture site.
- 5 2. A drug selected from the group consisting of at least one bisphosphonate when used for treating a fractured bone.
 - 3. The drug of claim 1 or claim 2 wherein the bisphosphonate is Zoledronate.
- 4. The drug of claim 1 or claim 2 wherein the drug is a combination of two or more bisphosphonates.
 - 5. The drug Zoledronate when used for promoting new bone formation.
 - 6. The drug of claim 5 when used for promoting new bone formation between a bone and a prosthesis, bone fixation device or any other bone or dental implant.
 - 7. The drug of any one of the preceding claims when administered to an individual as a single dose.
 - 8. The drug of any one of the preceding claims when administered to an individual perioperatively.
 - 9. Use of a drug selected from the group consisting of at least one bisphosphonate for the manufacture of a medicament for promoting new bone formation at a fracture site.
 - 10. Use of a drug selected from the group consisting of at least one bisphosphonate for the manufacture of a medicament for treating a fractured bone.
- 25 11. Use of the drug of claim 9 or claim 10 wherein the drug is Zoledronate.
 - 12. Use of the drug of claim 9 or claim 10 wherein the drug is a combination of two or more bisphosphonates.
 - 13. Use of the drug Zoledronate for the manufacture of a medicament for promoting new bone formation.
- 30 14. A method for treating a fractured bone, the method including administering to a subject with a fractured bone a therapeutically effective amount of a drug selected from the group consisting of at least one bisphosphonate.
 - 15. The method of claim 14 wherein the drug is administered to the subject as a single dose.
 - 16. The method of claim 15 wherein the single dose of drug is

AMENDED SHEET

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administered at an early stage of treatment of the fractured bone.

- 17. The method of claim 14 wherein the mode of administration is as a perioperative intravenous infusion.
- 18. The method of claim 14 wherein the mode of administration is oral.
- 19. The method of claim 14 wherein the mode of administration is transdermal.
 - 20. A method of treating a fractured bone, the method including the steps of:
 - (a) administering to a subject with a fractured bone a therapeutically effective amount of a drug selected from the group consisting of at least one bisphosphonate; and
 - (b) providing a vibratory stimulus to the fractured bone.
 - 21. The method of claim 20 wherein the vibratory stimulus is provided by ultrasound stimulation or vibration stimulation.
 - 22. The method of claim 20 or claim 21 wherein the vibratory stimulus includes periodically providing a vibratory stimulus at the resonant frequency of the bone.
 - 23. The method of claim 22 wherein the resonant frequency is calculated as a function of the bone's vibratory response to the vibratory stimulus.
 - 24. The method of any one of claims 20 to 23 wherein the vibratory stimulus is provided at a late stage in the treatment of the fractured bone.
 - 25. The method of any one of claims 20 to 23 wherein the step of providing a vibratory stimulus is concurrent with the step of administering a therapeutically effective amount of the drug.
- 25 26. The method of claim 25 wherein the vibratory stimulus is provided and the therapeutically effective amount of the drug is administered at an early stage in the treatment of a fractured bone.
 - 27. A drug selected from the group consisting of at least one bisphosphonate when used for promoting new bone formation at a fracture site in an individual suffering from delayed union of a fracture.
 - 28. A method for promoting new bone formation at a fracture site in a subject suffering from delayed union of a fracture, the method including administering to the subject a therapeutically effective amount of a drug selected from the group consisting of at least one bisphosphonate.
- The method of claim 28 wherein the at least one bisphosphonate is administered parenterally as a single dose at or near the time of surgery.



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- 30. The method of claim 29 wherein a further parenteral dose of the at least one bisphosphonate is administered about four to six weeks after the initial dose.
- 31. The method of claim 29 wherein further oral doses of the at least one bisphosphonate are administered in a daily or second daily regimen commencing about four to six weeks after the initial dose for a period of about two months or until sufficient new bone has been formed.
- 32. A method of promoting new bone formation in a subject, the method including the steps of surgically performing the procedure of distraction osteogenesis and administering to the subject a drug selected from the group consisting of at least one bisphosphonate.
- 33. The method of claim 32 wherein the at least one bisphosphonate is administered parenterally as a single dose at or near the time of surgery.
- 34. The method of claim 33 wherein a further parenteral dose of the at least one bisphosphonate is administered either at the end of the distraction period or up to three months after the initial dose.
- 35. The method of claim 33 wherein further oral doses of the at least one bisphosphonate are administered in a daily, second daily or weekly regimen.
- 36. The method of claim 35 wherein the regimen commences about one to three months after the initial parenteral dose for a period of about two months.